IN THE CLAIMS

1. - 14. (Cancelled)

15. (Currently Amended) A compound of formula (lb):

wherein R^a is selected from halogen, hydroxy, cyano, nitro, amino, hydroxycarbonyl, C_1 - C_6 alkyl, C_1 - C_6 alkenyl, C_1 - C_6 alkynyl, hydroxy C_1 - C_6 alkyl, C_1 - C_6 alkoxy C_1 - C_6 alkyl, perfluoro C_1 - C_6 alkoxy, C_1 - C_6 alkylamino, di- C_1 - C_6 alkylamino, amino C_1 - C_6 alkyl, C_1 - C_6 alkylamino C_1 - C_6 alkyl, C_1 - C_6 alkylamino C_1 - C_6 alkyl, C_1 - C_6 alkylamino, C_1 - C_6 alkylthio, C_1 - C_6 alkylamino, aminosulfonyl, C_1 - C_6 alkylaminosulfonyl, C_1 - C_1 - C_2 - C_3 - C_4 - C_4 - C_4 - C_5 - C_5 - C_6 - C_6 - C_6

 R^b is selected from hydrogen, halogen, hydroxy, (C_1-C_6) alkoxy cyano, nitro, amino, hydroxycarbonyl, C_1-C_6 alkyl, C_1-C_6 alkenyl, C_1-C_6 alkynyl, C_1-C_6 alkoxy, hydroxy C_1-C_6 alkyl, C_1-C_6 alkoxy C_1-C_6 alkyl, perfluoro C_1-C_6 alkyl, perfluoro C_1-C_6 alkylamino, diacute C_1-C_6 alkylamino, amino C_1-C_6 alkyl, C_1-C_6 alkylamino C_1-C_6 alkylamino C_1-C_6 alkylamino C_1-C_6 alkyl, C_1-C_6 alkylamino C_1-C_6 alkyl, C_1-C_6 alkylamino, C_1-C_6 alkylthio, C_1-C_6 alkylthiocarbonyl, C_1-C_6 alkylthioxo, C_1-C_6 alkylthioxo, C_1-C_6 alkylsulfonyl, C_1-C_6 alkylsulfonylamino, aminosulfonyl, C_1-C_6 alkylaminosulfonyl, di- C_1-C_6 alkylaminosulfonyl, $C_1-C_$

16. -17. (Cancelled)

18. (Currently Amended) A compound of formula (1b) according to claim 15 which is selected from the group consisting of:

(2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid;

(2S,4S)-4-(2,3-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; and

(2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid; or a pharmaceutically acceptable salt thereof.

19. (Cancelled)

- 20. (Previously Presented) A pharmaceutical composition comprising a compound of formula (1b) according to claim 15, or a pharmaceutically acceptable salt thereof, and one or more pharmaceutically acceptable excipients, diluents or carriers.
- 21. (Previously Presented) A pharmaceutical composition comprising a compound of formula (1b) according to claim 15, or a pharmaceutically acceptable salt thereof, and at least one other therapeutically active agent.
- 22. (Currently Amended) A <u>pharmaceutical composition combination</u> according to claim 21, wherein the other therapeutically active agent is a PDEV inhibitor selected from sildenafil, vardenafil, tadalafil, 1-{6-ethoxy-5-[3-ethyl-6,7-dihydro-2-(2-methoxyethyl)-7-oxo-2H-pyrazolo[4,3-d]pyrimidin-5-yl]-3-pyridylsulfonyl}-4-ethylpiperazine, 5-(5-acetyl-2-butoxy-3-pyridinyl)-3-ethyl-2-(1-ethyl-3-azetidinyl)-2,6-dihydro-7*H*-pyrazolo[4,3-d]pyrimidin-7-one and 5-[2-ethoxy-5-(4-ethylpiperazin-1-ylsulphonyl)pyridin-3-yl]-3-ethyl-2-[2-methoxyethyl]-2,6-dihydro-7*H*-pyrazolo[4,3-d]pyrimidin-7-one.

23. - 26. (Cancelled)

- 27. (Previously Presented) The compound (2*S*,4*S*)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid, or a pharmaceutically acceptable salt thereof.
- 28. (Withdrawn) The compound (2S,4S)-4-(2,3-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid, a pharmaceutically acceptable salt thereof.
- 29. (Withdrawn) The compound (2S,4S)-4-(2,5-Difluoro-benzyl)-pyrrolidine-2-carboxylic acid, or a pharmaceutically acceptable salt thereof.
- 30. (New) The salt, (2S,4S)-4-(3-Fluoro-benzyl)-pyrrolidine-2-carboxylic acid mono hydrochloride salt.